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REMARKS

I. Status Summary

Claims 1-19 are pending in the present application. Election by applicants of Group I, claims 1-17 has been acknowledged by the U.S. Patent and Trademark Office (hereinafter "the Patent Office") and claims 1-3, 5-10 and 12-17 currently examined. Claims 18-19 have been withdrawn from prosecution at this time and claims 4 and 11 have been canceled. Claims 1, 5-6, 8-9, and 13-16 are currently amended.

Claims 1, 2, 9 and 16-17 presently stand rejected under 35 USC § 103(a) as allegedly being obvious over U.S. Patent No. 5,252,466 to Cronan (hereinafter "Cronan") in view of U.S. Patent No. 5,283,173 to Fields et al. (hereinafter "Fields et al."). Claims 1, 2, 5-9 and 12-16 presently stand rejected under 35 USC § 103(a) as allegedly being obvious over Cronan in view of Fields et al. and further in view of the journal article to Rigaut et al. (Nature Biotech., 17: 1030-1032, 1999; hereinafter "Rigaut et al."). Claims 1-3, 5-10 and 12-17 presently stand rejected under 35 USC § 103(a) as allegedly being obvious over Cronan in view of Fields et al., further in view of Rigaut et al., and further in view of U.S. Patent No. 6,114,111 to Luo et al. (hereinafter "Luo et al.").

II. Response to the Rejections Under 35 U.S.C. § 103

II.A. Cronan in view of Fields et al.

Claims 1, 2, 9 and 16-17 have been rejected under 35 U.S.C. § 103(a) as being unpatentable over <u>Cronan</u> in view of <u>Fields et al.</u>

Applicants respectfully submit that one of ordinary skill in the art would not have had a motivation to combine <u>Cronan</u> with the teachings of <u>Fields et al.</u> and, in any case, even if the references were to be combined the combination would fail to teach or suggest every element of claims 1, 2, 9 and 16-17. <u>Cronan</u> does not teach "binding partners" to a protein of interest as the phrase "binding partners" is used by

Applicants and, accordingly, one of ordinary skill in the art could not have had a motivation to combine <u>Cronan</u> and <u>Fields et al.</u> on such a basis. The Examiner seems to be suggesting that <u>Cronan</u> does teach obtaining such "binding partners", and the reference is only lacking in teaching the identification of the binding partners. (See Official Action, page 4, first full paragraph). To the contrary, the binding factors in <u>Cronan</u> are *provided* as part of the disclosed procedure for purifying the protein of interest and, thus, there is no need to "identify" these binding partners.

It is respectfully submitted that there is a misapprehension of the meaning of "binding factors" as used in the <u>Cronan</u> reference versus use of the phrase in Applicants' disclosed subject matter. <u>Cronan</u> is using "binding partners" to mean the binding partner for the molecule which is attached by post-translational modification to the engineered site. This is why the <u>Cronan</u> reference specifically states: "providing a binding partner that binds to the fusion protein only after it has been modified". (See, e.g., the Abstract at page 1; the Summary at column 5, line 68 (emphasis added)). <u>Cronan</u> is directed to protein purification by providing a known binding partner to the tag as part of the purification procedure. In contrast, Applicants are using "binding partners" to mean molecules that bind to the protein of interest at areas other than the post-translational modification. The binding partners disclosed by Applicants are not provided, and are not part of the procedure for separating the protein of interest from the cellular extract as is the case in <u>Cronan</u>.

In the claims, it does not matter whether the "binding partner" binds before or after the occurrence of the post-translational modification. Applicants' binding partner is not binding to the molecule added by post-translational modification. For example, each of amended independent claims 1, 9 and 16 is directed to obtaining or screening for "binding partners", and each of these claims includes a step for contacting the tagged fusion protein with an affinity purification reagent while the tagged fusion protein is present in the cellular extract, and each claim contains a step for identifying binding partners that bind to the protein of interest. Applicants describe binding of both an affinity purification reagent and binding partners to the protein of interest. The binding partners described in Cronan at best relate to the

"affinity purification reagent", rather than the "binding partners" in the claims. Claims 1 and 9 are currently amended at step (b) to further point out and clarify the difference between the binding partners to be identified and the affinity purification reagent used in separating the fusion protein from the cellular extract. Support for the amendment to claims 1 and 9 can be found throughout the specification as filed, including, in particular, the Summary at page 4, lines 1-8; the Detailed Description at page 9, lines 5-15 and 24-31; page 14, lines 22-24; page 17, lines 12-20; page 18, lines 16-19; page 22, lines 5-21; and pages 24-25. No new matter has been added.

The phrase "binding partner" is used by both <u>Cronan</u> and Applicants. However, the phrase is used differently in each case. This difference in meaning would have been readily apparent to one of ordinary skill in the art at the time the instant application was filed. Accordingly, the skilled artisan would not have had a motivation to combine <u>Cronan</u> with <u>Fields et al.</u> There must be some suggestion or motivation, either in the references themselves or in the knowledge generally available to one of ordinary skill in the art, to modify the reference or to combine reference teachings. *In re Vaec*k, 947 F.2d 488, 20 USPQ2d 1438 (Fed. Cir. 1991). Prior art references must be considered in their entirety, i.e., as a <u>whole</u>, including portions that would lead away from the claimed invention. *W.L. Gore & Associates, Inc. v. Garlock, Inc.*, 721 F.2d 1540, 220 USPQ 303 (Fed. Cir. 1983), cert. denied, 469 U.S. 851 (1984).

A skilled artisan would not have had a motivation based on use of the phrase "binding factors" in <u>Cronan</u> to combine the reference with the teachings of <u>Fields et al.</u> While <u>Fields et al.</u> is directed to the identification of protein-protein interactions within a cell, <u>Cronan</u> only teaches separation of a protein from a cellular extract by provision of a known binding partner/affinity purification reagent. Since <u>Cronan</u> does not teach or suggest identifying non-affinity purification reagent binding partners to the protein of interest, one of ordinary skill in the art desiring to identify such binding partners would not be motivated to turn to <u>Cronan</u> to look for desired components, as <u>Cronan</u> does not teach or suggest a key component of the endeavor, that is, identifying binding partners of a protein of interest. As such, Applicants respectfully

submit there is no suggestion or motivation to combine the teachings of <u>Cronan</u> and <u>Fields et al.</u>

Furthermore, even if Cronan were combined with Fields et al., every element of claims 1, 2, 9 and 16-17 could not be present without destroying operability. To establish prima facie obviousness, each and every claim limitation must be taught or suggested by the prior art. In re Royka, 490 F.2d 981, 180 USPQ 580 (CCPA 1974). Fields et al. use a yeast two-hybrid system to identify the interaction between two proteins of interest. In Fields et al., the protein-protein interaction is identified by reconstitution of transcriptional activation of a reporter gene. Once the Cronan and Fields et al. methods are combined, it is not possible to perform every step of independent Claims 1, 9 and 16. For example, part (d) of claims 1, 9 and 16 describes separating the tagged fusion protein-affinity purification reagent complex from the cell extract, and part (e) recites identifying any binding partners of the protein of interest. However, in the <u>Fields et al.</u> method, separation of one or both of " the proteins of interest from the cell extract would make detection of the binding partner impossible, because the transcriptional activation reporter system requires the presence of the cellular extract to function. When the methods of Cronan and Fields et al. are combined, every element of the claims cannot be present without destroying operability. Therefore, the combination of Cronan and Fields et al. fails to teach or suggest every element of independent claims 1, 9 and 16. Accordingly, the claims cannot be rendered obvious by the proposed combination.

Applicants respectfully submit that because there is no motivation to combine the teachings of <u>Cronan</u> and <u>Fields et al.</u>, and the proposed combination would result in the inoperability of independent claims 1, 9 and 16, the cited combination does not support the instant rejection of these claims. Claims 2 and 17 depend from independent claims 1 and 16, respectively and, therefore, support for the rejection of these claims is similarly lacking. Hence, Applicants respectfully submit that a *prima facie* case of obviousness has not been established, and Applicants respectfully request the instant rejection of claims 1, 2, 9 and 16-17 under 35 U.S.C. § 103(a) be withdrawn. Allowance of these claims is also respectfully requested,

II.B. <u>Cronan</u> in view of <u>Fields et al.</u>, in further view of <u>Rigaut et a</u>l.

Claims 1, 2, 5-9 and 12-16 have been rejected under 35 U.S.C. § 103(a) as being unpatentable over <u>Cronan</u> in view of <u>Fields et al.</u>, and in further view of <u>Rigaut</u> et al.

Applicants respectfully submit that one of ordinary skill in the art would not have had any motivation to combine <u>Cronan</u> with the teachings of either <u>Fields et al.</u> or <u>Rigaut et al.</u> and, in any case, even if the references were to be combined, the combination would fail to disclose every element of claims 1, 2, 5-9 and 12-16. As described above, one of ordinary skill in the art would not have been motivated to combine the <u>Cronan</u> and <u>Fields et al.</u> references, as there is no suggestion or teaching in <u>Cronan</u> to identify non-affinity reagent-type binding partners to a protein of interest. Furthermore, the proposed combination renders impossible the performance of every element of independent claims 1, 9 and 16 without destroying operability. The addition of the <u>Rigaut et al.</u> reference fails to rectify this deficiency and, in any case, <u>Rigaut et al.</u> teaches away from combination with <u>Cronan</u>.

One of ordinary skill in the art would not have been motivated to combine Rigaut et al. with Cronan, as Rigaut et al. teaches away from Cronan. Furthermore, even if Rigaut et al. were combined as proposed, the resulting combination would still fail to teach or suggest every element of amended claims 1, 2, 5-9 and 12-16. Rigaut et al. teaches transforming a yeast cell with a fusion protein comprising a heterologous protein with two affinity tags (two IgG-binding units of protein A (ProtA) and a calmodulin binding peptide (CBP)) and a TEV cleavage site. Rigaut et al. designate the fusion cassette encoding the CBP, TEV cleavage site, and ProtA the tandem affinity purification (TAP) tag due to its consisting of two different affinity tags. Rigaut et al. describe a method for protein purification and protein complex characterization using the serial two high-affinity tag purification system. Rigaut et al. teaches away from Cronan by expressly stating and demonstrating the necessity of having the two-step serial affinity purification procedure. (See Rigaut et al. at page 1031, column 1).

Independent claims 1, 9 and 16 presently recite a cell transformed to express a fusion protein having a single post-translational modification sequence resulting in a singly tagged fusion protein. Support for the claim amendments can be found throughout the specification and claims as filed and in particular at the Summary, page 4, lines 1-8; Figure 1; and Example 1, pages 19-26. No new matter has been added. Given the express requirement for using tandem two affinity tag purification in Rigaut et al., one of ordinary skill in the art would not have had a reasonable expectation of success in combining Rigaut et al. with the method of Cronan, to obtain the single affinity reagent separation procedure as recited in independent claims 1, 9 and 16. Indeed, Rigaut et al. is believed to provide evidence that the success of the instantly claimed methods was contrary to the expectations of skilled artisans at the time of filing of the present U.S. patent application. Accordingly, one of ordinary skill in the art would not have expected success in the proposed. combination, and consequently would not have been motivated to combine Cronan with <u>Rigaut et al.</u> in such manner.

Furthermore, even if Rigaut et al. were combined with Cronan and Fields et al., every element of claims 1, 2, 5-9 and 12-16 would not be taught or suggested. To establish prima facie obviousness, each and every claim limitation must be taught or suggested by the prior art. In re Royka, 490 F.2d 981, 180 USPQ 580 (CCPA 1974). As described above, the Rigaut et al. reference teaches away from the claim limitations of a single post-translational modification sequence and a singly tagged fusion protein. Accordingly, the combination of Cronan, Fields et al. and Rigaut et al. fails to teach or suggest every element of the claims. Therefore, claims 1, 9 and 16 cannot be rendered obvious by the proposed combination.

Applicants respectfully submit that because a skilled artisan could not have expected success in combining Cronan and Fields et al. and the proposed combination of Cronan, Fields et al. and Rigaut et al. would, in any case, fail to teach or suggest every element of independent claims 1 and 9 and 16 as the claims would be rendered inoperable, the cited combination does not support the instant rejection of these claims. Claims 2, 5-8 and 12-15 depend from independent claims 1 and 9,

and therefore support for the rejection of these claims is similarly lacking. Hence, Applicants respectfully submit that a *prima facie* case of obviousness has not been established, and Applicants respectfully request the instant rejection of claims 1, 2, 5-9 and 12-16 under 35 U.S.C. § 103(a) be withdrawn. Allowance of these claims is also respectfully requested.

II.C. <u>Cronan</u> in view of <u>Fields et al.</u>, in view of <u>Rigaut et al.</u> and in view of <u>Luo</u> et al.

Claims 1-3, 5-10 and 12-17 have been rejected under 35 U.S.C. § 103(a) as being unpatentable over <u>Cronan</u> in view of <u>Rigaut et al.</u>, in further view of <u>Fields et al.</u>, and in further view of <u>Luo et al.</u>

Applicants respectfully submit that one of ordinary skill in the art would not have had any motivation to combine <u>Cronan</u> with the teachings of either <u>Fields et al.</u> or <u>Rigaut et al.</u> or <u>Luo et al.</u> and, in any case, even if the references were to be combined, the combination would fail to teach or suggest every element of claims 1-3, 5-10 and 12-17 and would render the methods inoperable. As described above, one of ordinary skill in the art would not have been motivated to combine <u>Cronan</u> with <u>Fields et al.</u> or <u>Rigaut et al.</u>, as there is no suggestion or teaching for the combination and, in any case, the combination would render impossible the performance of every element of independent claims 1, 9 and 16 without destroying operability. The addition of the <u>Luo et al.</u> reference fails to rectify this deficiency, and in a manner similar to the <u>Fields et al.</u> reference, would render the claims inoperable.

Similar to the case of <u>Fields et al.</u> described above, one of ordinary skill in the art would not have had a motivation based on use of the phrase "binding factors" in <u>Cronan</u> to combine the reference with the teachings of <u>Luo et al.</u> While <u>Luo et al.</u> is directed to the identification of protein-protein interactions within a cell, <u>Cronan</u> only teaches separation of a protein from a cellular extract by provision of a known binding partner/affinity purification reagent. Since <u>Cronan</u> does not teach or suggest identifying non-affinity purification reagent binding partners to the protein of interest, one of ordinary skill in the art desiring to identify such binding partners would not be

motivated to turn to Cronan to look for desired components, as Cronan does not teach or suggest a key component of the endeavor, that is, identifying binding partners of a protein of interest. As such, applicants respectfully submit there is no suggestion or motivation to combine the teachings of Cronan and Luo et al.

Furthermore, even if Cronan were combined with Luo et al., every element of claims 1-3, 5-10 and 12-17 could not be present without destroying operability. To establish prima facie obviousness, each and every claim limitation must be taught or suggested by the prior art. In re Royka, 490 F.2d 981, 180 USPQ 580 (CCPA 1974). In Luo et al., protein-protein interactions are identified by reconstitution of transcriptional activation of a reporter gene. Once the Cronan and Luo et al. methods are combined, it is not possible to perform every step of independent Claims 1, 9 and 16. For example, part (d) of claims 1, 9 and 16 describes separating the singly tagged fusion protein-affinity purification reagent complex from the cell extract, and part (e) recites identifying any binding partners of the protein of interest. However, in the <u>Luo et al.</u> method, separation of one or both of the proteins of interest from the cell extract would make detection of the binding partner impossible. because the transcriptional activation reporter system requires the presence of the cellular extract to function. When the methods of Cronan and Luo et al. are combined, every element of the claims cannot be present without destroying operability. Therefore, the combination of Cronan and Luo et al. fails to teach or suggest every element of independent claims 1, 9 and 16. Accordingly, the claims cannot be rendered obvious by the proposed combination.

Applicants respectfully submit that because a skilled artisan would not have been motivated to combine <u>Cron</u>an and Luo<u>et al.</u> and the proposed combination of Cronan, Fields et al., Rigaut et al. and Luo et al. would, in any case, render inoperable independent claims 1, 9 and 16, the cited combination does not support the instant rejection of these claims. Claims 2-3, 5-8, 10, 12-15 and 17 depend from independent claims 1, 9 and 16 and, therefore, support for the rejection of these claims is similarly lacking. Hence, Applicants respectfully submit that a prima facie case of obviousness has not been established, and Applicants respectfully request

the instant rejection of claims 1-3, 5-10 and 12-17 under 35 U.S.C. § 103(a) be withdrawn. Allowance of these claims is also respectfully requested.

CONCLUSION

In light of the above amendments and remarks, it is respectfully submitted that the present application is now in proper condition for allowance, and an early notice to such effect is earnestly solicited.

If any small matter should remain outstanding after the Patent Examiner has had an opportunity to review the above Remarks, the Patent Examiner is respectfully requested to telephone the undersigned patent attorney in order to resolve these matters and avoid the issuance of another Official Action.

DEPOSIT ACCOUNT

Although it is believed that no fee is due, the Commissioner is hereby authorized to charge any fees associated with the filling of this Response, or credit any overpayment, to Deposit Account No. <u>50-0426</u>.

Respectfully submitted,

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